Message

From: Overstreet, Anne [overstreet.anne@epa.gov]
Sent: 5/7/2020 6:32:21 PM
To: Mendelsohn, Mike [Mendelsohn.Mike@epa.gov]; McNally, Robert [Mcnally.Robert@epa.gov]
Subject: RE: Draft email to Keith

Here is my concern and a couple of questions: what process do we use on the back end to ensure that what Oxitec is claiming as CBI is consistent with how the regs and guidance docs define it? I'm concerned that Oxitec will strike information that does not suit them from our risk assessment that is not, in fact, confidential. Does ITRMD verify their specific redacted portions? Would this circumvent the 6 to 8 month process that OGC noted or is this the first step in that process?

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http://www.epa.gov/pesticides

From: Mendelsohn, Mike <Mendelsohn.Mike@epa.gov>
Sent: Thursday, May 07, 2020 2:06 PM
To: McNally, Robert <Mcnally.Robert@epa.gov>; Overstreet, Anne <overstreet.anne@epa.gov>
Subject: RE: Draft email to Keith

Anne and Bob,

Earl is sending an email to Keith, but before I provide the details he needs, can you weigh in on the statement

Ex. 5 Deliberative Process (DP)

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Thanks.

Mike

From: Sadowsky, Don <Sadowsky.Don@epa.gov>
Sent: Thursday, May 07, 2020 1:37 PM
To: Mendelsohn, Mike <Mendelsohn.Mike@epa.gov>; Ingram, Earl <Ingram.Earl@epa.gov>
Cc: Kaczmarek, Chris <Kaczmarek.Chris@epa.gov>; Schwarz, Stephanie <Schwarz.Stephanie@epa.gov>
Subject: Draft email to Keith

Mike and Earl --

Here's the guts of an email or letter I suggested that Earl send to Keith, with Mike filling in the blanks.

> EPA intends to place in the public record the following documents related to **EPA's issuance of Oxitec's Experimental Use Permit 93167-EUP-2:**
>
> **Human Health and Environmental Risk Assessment for the New Product OX5034 dated April 30, 2020**

Ex. 5 Deliberative Process (DP)

Does this work for you?

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